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METHOD FOR ELIMINATING DEFECTS IN SILICON ELEMENTS THROUGH SELECTIVE ETCHING

The present invention relates to a method for eliminating eruptions, impurities, and/or damage in the crystal lattice by selectively etching silicon elements, in particular of surface-plated parts of a silicon wafer, according to the definition of species in Claim 1.

BACKGROUND INFOR WHITEON

A known method for producing silicon power diodes is to cut a number of individual diodes from a silicon wafer that has been doped on both sides with an n- or p-type dopant in full-surface doping steps to generate a pn junction and subsequently plated all over on both sides, by first sawing the entire wafer into squares or hexagons measuring roughly 5 x 5 mm² and then suitably building up and electrically bonding such individual silicon surface diodes. The silicon wafer is doped and plated in a manner that is known per se, with the surface plating layer being made, for example, of a chromium-nickel-vanadium-silver alloy (CrNiVAg).

The sawing step, which is necessary to produce individual may results in irregular diodes from the silicon wafer, efter results in irregular eruptions on the sawn edge or impurities and damage in the crystal lattice so that the pn junctions in the sawn-out individual diodes are partially or regionally damaged. This damage then leads to higher leakage currents, which, in the end, renders the created diode unusable at this stage.

Therefore, in known methods for producing silicon power diodes of this type, the sawing step must be followed by a wetchemical overetching of the diodes, thus stripping the damaged silicon areas and restoring a suitably intact crystal lattice in the pn junction region.

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The chemical etch-stripping of the damaged silicon areas to them followed by passivation of the exposed silicon edge to protect the pn junctions against environmental influences and to reliably prevent the electronic properties from again deteriorating over the life of the diode. The latter object is considered to be achieved by the related art.

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However, there are at present no satisfactory means of etchstripping the damaged silicon areas at the sawn edge. The wetchemical etching method frequently used today results, for example, in unfavorable etching profiles with greater risk of electrical breakdown, due to its pronounced doping selectivity, as well as in disadvantageous yields, due to occasional rejects during etching. In addition, the etching profile produced by wet-chemical etching also reduces the diode's mechanical stability.

Furthermore, according to the currently known methods, only the fully mounted diodes are exposed to the aqueous etching solution, which involves expensive handling of the individual diodes. To summarize, therefore, the methods known from the related art for overetching silicon elements sawn out of a silicon wafer are described by the following process steps:

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Wet-chemical etching of the sawn-out silicon elements, i.e., diodes, mounting of the diodes, immersion of the mounted diodes in etching baskets into etching basins, neutralization of the etching solution, and subsequent thorough rinsing and drying of the diodes.

The agents needed for the known method are an etching solution, a neutralization solution, and hydrogen peroxide, resulting in serious environmental pollution due to the materials used as well as high energy consumption and the use of de-ionized water. Thus, wet-etching takes place, for

needed for subsequent cleaning of the etched, mounted diodes.

SUMMALL OF THE INVENTION
Advantages of the Invention

The method according to the present invention having the characterizing features of the main claim has the advantage over the related art that it can be used to etch silicon elements, in particular surface-plated, sawn-out parts of a silicon wafer, using a gaseous etching medium that selectively etches almost exclusively silicon through a chemical reaction, thus producing gaseous reaction products. During this selective etching process, surface eruptions and/or impurities, and/or damage in the crystal lattice of the silicon element, in particular, are eliminated at the same exemplary method time. The method according to the present invention is further for advantageously suitable for stripping damaged silicon zones, in particular of sawn-out silicon elements, like those that occur, for example, when producing silicon power diodes. It is reliable, economical, and overcomes the known disadvantages of fluid etching media.

example, at temperatures above 90°C, and a rinsing cascade is

One particular advantage is that the method according present invention is a "batch process", i.e., a process on the wafer level. Thus, one particular advantage is that the individual chips do not need to be handled, which means that all sawn-out silicon elements, i.e., chips, of a wafer can be etched simultaneously, which saves a great deal of space and requires only one process step, i.e., handling step. necessary, the method according to the present invention can also be used to overetch fully mounted individual diodes, as is currently the case. In doing this, it is necessary only to replace the wet-etching process that is known per se with an etching process using a gaseous etching medium.

exemplary method A further advantage is that the method according to the present invention for selective etching does not use a fluid,

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elements that have been, in particular, sawn out. Moreover, exemplay we had the present invention is also less selective toward doping concentrations, resulting in advantageous etched edge profiles and, in particular, prevents a "boron balcony" from forming on the sawn-out and overetched diodes, at the same time increasing mechanical stability and reducing failure rates.

thus achieving a clean gas-phase etching of the silicon

is believed to be the selective removal of damaged Another great advantage is silicon zones, eruptions or impurities, as well as the <u>EXEMPLAN</u> Method planarizing effect of the metho according to the present invention. Due to the high selectivity of the gaseous etching medium used and the chemical reaction that this produces on the surface of the silicon element, and particularly due to the selectivity of this reaction on damage in these zones, it is possible to etch-strip significantly more material, with damaged areas automatically undergoing more aggressive etching, particularly in the edge areas of the sawn-out silicon elements.

Advantageous embodiments of the present invention are provided by the features mentioned in the subclaims:

exemplant method 25 Thus, the method according to the present invention offers the advantage that the etching rates can be selectively adjusted, for example via the composition of the gaseous etching medium. This makes it possible to set, if necessary, low etchstripping rates, thereby increasing reproducibility and thus , reducing overall process time. However, it is also possible to 30 set very high etching rates, particularly when using chlorine trifluoride or bromine trifluoride, and thus significantly shorten the actual etching time.

The very high selectivity of the 35 present invention toward non-silicon materials is due to a surface-catalytic initiation of the etching reaction, so that

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a large number of materials are also advantageously suitable for use as the etching mask.

For example, even a surface-plating of the silicon wafer, which serves as the later diode plating, can be used as the etching mask while etching with the gaseous etching medium without this plating itself being significantly corroded. Another advantage in this regard is that a sawing sheet used and an adhesion layer provided between this sawing sheet and the silicon wafer attached to it are also not significantly corroded, due to the high selectivity of the etching medium toward non-silicon materials. Furthermore, an additional lacquer layer can be advantageously applied all over in a manner that is known per se, for example by spin-coating, thus further protecting the surface plating of the silicon wafer.

In the case of the interhalogen or fluorine-noble gas compounds suitable for the method according to the present invention, suitable process conditions further ensure that no free chlorine, bromine or iodine occurs, leaving only a very low risk of corrosion after completion of the etching process.

Further advantages of the method according to the present invention include low energy consumption because there is no need to heat, for example, the diodes or etching baths, low chemical consumption, and environmental compatibility. For example, any waste gases produced can be very easily disposed of without harmful effects through after-treatment, eliminating hazardous waste. An advantageously suitable method includes without gases in lime water, i.e., an aqueous calcium hydroxide solution in a "gas washer" (Ca(OH)₂ + 2HF -> CaF₂ + 2 H₂O).

Chlorine trifluoride, bromine trifluoride, iodine pentafluoride, or xenon difluoride that are in a gaseous state or have been converted to the gaseous phase, or a mixture of

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these compounds, are particularly suitable as the etching medium. In this case, silicon tetrafluoride, for example, is produced as the reaction product.

DETAILED DESCRIPTION

5 Description of Embodiments

The method according to the present invention for eliminating eruptions, impurities, or damage in the crystal lattice of silicon elements by selective etching of silicon generally makes use of the characteristic of certain fluorine compounds, known as interhalogens or fluorine-noble gas compounds, to spontaneously etch silicon, i.e., through contact between the fluorine compound and silicon, where this etching action takes place from the gaseous phase, and gaseous reaction products are produced. To do this, a gaseous silicon-etching fluorine compound, for example, is supplied to a reaction chamber of a supplied to a suppli be etched were previously placed. Adsorption of the gas molecules of the gaseous etching medium on the accessible silicon surfaces results in spontaneous, surface-catalytic fragmentation of the etching medium used, thus releasing fluorine radicals that react with the silicon to form a volatile product, for example SiF_x (x = 2, 3, 4). The best known product of a reaction of this type is, for example, "stable silicon tetrafluoride SiF4. The mechanism, which is known per se, can be described as follows, where X is Cl, Br, I or Xe, and n is the number of fluorine atoms in the particular compound:

30 1. Adsorption:

$$XF_n \rightarrow XF_n^{adsorb}$$

- 2. Surface-catalytic decomposition: $XF_n^{adsorb} o XF_{n-1}^{adsorb} + F^{*,adsorb}$
- 3. Chemical transformation: $Si + xF^{*,adsorb} \rightarrow SiF_{r}$ (x = 1, 2, 3, 4)

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4. Desorption of reaction products: $SiF_x \rightarrow SiF_x \uparrow$ (x = 2, 3, 4)

in particular: $Si + 4F^{*adsorb} \rightarrow siF_4 \uparrow$

Examples of suitable compounds of the XF_n type are the interhalogen compounds chlorine trifluoride, bromine trifluoride and iodine pentafluoride, as well as the noble gas fluoride xenon difluoride. The chlorine trifluoride and available bromine trifluoride compounds are known per se in the semiconductor industry, where they are commonly used to clean wafers or wafer handling devices.

While chlorine trifluoride and bromine trifluoride are liquids with a vapor pressure of 1 bar at 150°C, iodine pentafluoride is a semiliquid, and xenon difluoride is a solid with a vapor pressure of approximately 20 mbar and 2 mbar, respectively, at 15°C. Thus chorine trifluoride and bromine trifluoride can be taken directly from a gas bottle, due to their high vapor pressure, while iodine pentafluoride and xenon difluoride must first be converted to the gaseous phase, using a suitable vaporizing apparatus, preferably at an elevated temperature. When using the liquid iodine pentafluoride, it is preferable to use a "bubbler", in which the liquid is "bubbled" with an inert gas, for example helium, as the carrier gas, and the gas mixture is then supplied to the reaction chamber.

In the case of the interhalogen compounds chlorine trifluoride and bromine trifluoride, stripping takes place in a first reaction step of the etching reaction, in which fluorine radicals are released to form stabile chorine fluoride (ClF) and unstable bromine fluoride (BrF), respectively, with two fluorine radicals being generated in each case.

In the case of iodine pentafluoride, the medium is first converted to the relatively stable iodine trifluoride, and in the case of the noble gas fluoride xenon difluoride, elementary xenon is formed in addition to the two fluorine radicals.

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When carrying out the method according to the present invention, one embodiment begins with a silicon wafer from which silicon power diodes are to be produced, with one side of the wafer being doped with an n-type dopant in a manner that is known per se, using an all-over doping step, and the other side being doped with a p-type dopant in a manner that is known per se, using an all-over doping step. A pn-junction thus forms over the entire interior of the silicon wafer.

Afterwards, the wafer that has been pretreated in this manner using an available is provided on both sides with an all-over CrnivAg plating that is known per se.

This silicon wafer is then attached to a commercially Male like Sawing Sheet Made available, flexible sawing sheet, which is known per se, made for example from polyvinyl chloride (PVC) or polycarbonate, using an adhesion layer provided on it, and sawn using a sawing method in a manner that is known per se into silicon elements measuring approximately 5 x 5 mm² and shaped like squares or hexagons, with the silicon elements being used as silicon power diodes at the end of the manufacturing process.

After the wafer has been sawn, the sawn-out silicon elements according to the present invention, while still attached to the sawing sheet, which means that no individual chips are yet produced at this stage, but only an entire unit of sawn-out silicon elements.

To ensure mechanical stability, the edge of the sawing sheet may be is preferably clamped in a fixed frame during sawing and during the course of the remaining process steps, so that it can be gripped with particular ease and treated automatically. After the wafer has been sawn, the sawing sheet is preferably first expanded to increase the distance between the silicon elements and thus give the gaseous etching medium used later

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on, for example chorine trifluoride, better access to the side walls of the individual sawn-out silicon elements. An expansion frame is preferably used for this purpose, forming a stable frame for handling the otherwise flexible sawing sheet.

In any case, it is important to carefully dry the sawn-out silicon elements after sawing the silicon wafer and expanding the sawing sheet, but before etching. This ensures that no moisture enters the reaction chamber of the reactor used later to carry out the actual selective etching of the silicon elements.

In doing this, it is advisable to load the sawn-out silicon elements that have been placed on the sawing sheet and are joined to the wafer into the reaction chamber of the reactor using a loading device, such as a load lock, which includes an evacuation and heating function, for example a radiation heater using corresponding lamps.

Pumping out the loading device to create a vacuum and simultaneously heating the wafer that was sawn into silicon elements and is located on the sawing sheet in the loading device, for example using a radiation heater, removes remnants of moisture particularly efficiently before the wafer that was sawn into silicon elements enters the actual reaction chamber of the reactor, where the described etching reaction takes place after introducing the gaseous etching medium. The presence of moisture greatly encourages corrosion effects at this stage and is therefore undesirable.

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To release fluorine radicals, the etching gases used require surface-catalytic decomposition that takes place only in connection with the actual etching reaction with the silicon, which means that the etching reaction is extraordinarily highly selective with respect to non-silicon materials. As a result, the etching conditions can be very easily set, for example by selecting a gaseous etching medium and

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concentration so that only the silicon surfaces, but not metals or plastics, are significantly corroded by the etching chemicals used.

Thus metal layers, in particular applied surface plating, as well as the sawing sheet used fully satisfy the requirements that an etching reaction mask must meet to strip only a sawn edge damaged during sawing, but not the entire silicon surface. If additional protection of the surface plating or metal layers against even minor etching corrosion is required, this can be easily accomplished by additionally applying, in particular spin coating, a lacquer layer that is known per se onto preferably the entire surface of the silicon wafer.

After the sawn silicon wafer, which has been attached to the sawing sheet and dried, enters the reaction chamber of the reactor, the gaseous etching medium is then introduced into this chamber. When using the interhalogen compounds chlorine trifluoride or bromine trifluoride, the medium is introduced through flow regulators or throttle valves that are known per se, where the process pressure can range from a low-pressure range, i.e., vacuum, to an atmospheric pressure range. Both of the gases mentioned can be used, for example, in conjunction with a process in the pressure range from 0.1 to 1,000 mbar under controlled flow conditions.

If the process is to take place at higher pressures, it is advantageous to dilute the reactive gas with an inert gas, for example helium. Diluting the gas with helium by a factor of 10 to 100 further makes it possible to very easily control the etching reaction and the etching rate and reduce the aggressiveness of the gaseous etching medium used, which also reduces the potential chemical corrosion of gas lines and the interior of the reaction chamber.

By using a diluted inert gas, it is further possible to work even at atmospheric pressure by limiting the partial pressure,

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for example of chlorine trifluoride or bromine trifluoride, to a range of some 10 mbar by correspondingly diluting these gases with helium, which is still sufficient for maximum etching rates, yet also allows the etching process to be efficiently controlled through the inflow of the etching species.

When using the noble gas fluoride xenon difluoride, the latter must first be thermally sublimed from a solid source. In this case, the process pressure when etching the sawn-out silicon elements is thus limited to the vapor pressure of xenon difluoride at the selected working temperature, for example 2 mbar at 20°C.

If iodine pentafluoride is used as the etching medium, the latter must first be converted to the gaseous phase in an evaporator, which limits the working pressure to approximately 20 mbar. Alternatively, however, one can also use a "bubbler" to "bubble" the liquid iodine pentafluoride with an inert gas, for example helium, and supply the gas mixture diluted in this manner to the reaction chamber. Both procedures are known per se to those skilled in the art.

After introducing the gaseous etching medium into the reaction chamber, the actual etching of the sawn-out silicon elements begins, where specifically the damaged areas in the crystalline structure, known as damage zones, preferably on the sawn edge are stripped, and the sawn surfaces are planarized.

The gaseous reaction products produced from the reaction between the gaseous etching medium used and the silicon surfaces are either pumped away continuously, if a throughflow system is used, or accumulate in a reaction chamber that is filled once and then closed, until all gaseous substances are finally pumped out of the reaction chamber upon completion of silicon element etching.

In any case, pumping must continue for a sufficiently long period, and the discharge pressure during pumping must be low enough, preferably less than 0.1 μ bar, to ensure that the reaction chamber no longer contains any leftover etching gas before the silicon elements sawn out of the silicon wafer and located on the sawing sheet are discharged from the reaction chamber of the reactor either through the load lock or via the loading device.

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When discharging the elements, it is further suitable to also provide a vacuum in the load lock between the reaction chamber and the surrounding atmosphere. This vacuum preferably has a pressure of less than 0.1 μ bar, which can be easily achieved by using a turbomolecular pump.

In doing this, the wafer that has been sawn into silicon elements and is attached to the sawing sheet is preferably reheated in the load lock after etching, for example using a radiation heater. This removes as much leftover etching species as possible before the wafer is removed from the load lock, i.e., leftover gaseous etching medium or leftover gaseous reaction products from the sawn-out, overetched silicon elements or from their surfaces affected by etching.

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Leftover media of this type remaining on the surface of the silicon element would result in corrosion when exposed to air, due to the effect of moisture in the air.

After the processed silicon elements have been discharged from the reactor, they are finally removed individually from the sawing sheet and built up into diodes in a manner that is known per se. The silicon edges are then passivated in a manner that is also known per se to ensure the stability of

the pn junctions and the high performance of the silicon power diodes over their life cycles.

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